

***Title : SPECT-OPT multimodal imaging enables accurate evaluation of radiotracers for  $\beta$ -cell mass assessments***

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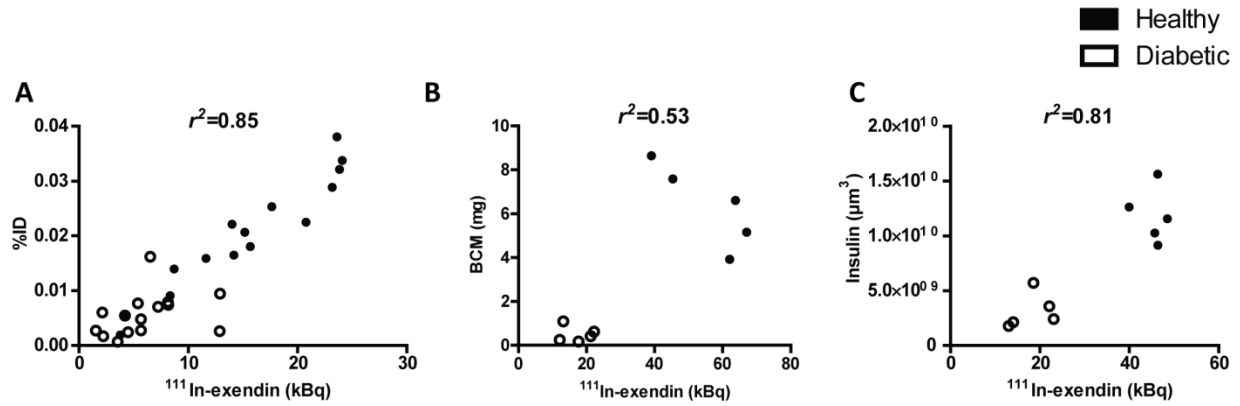
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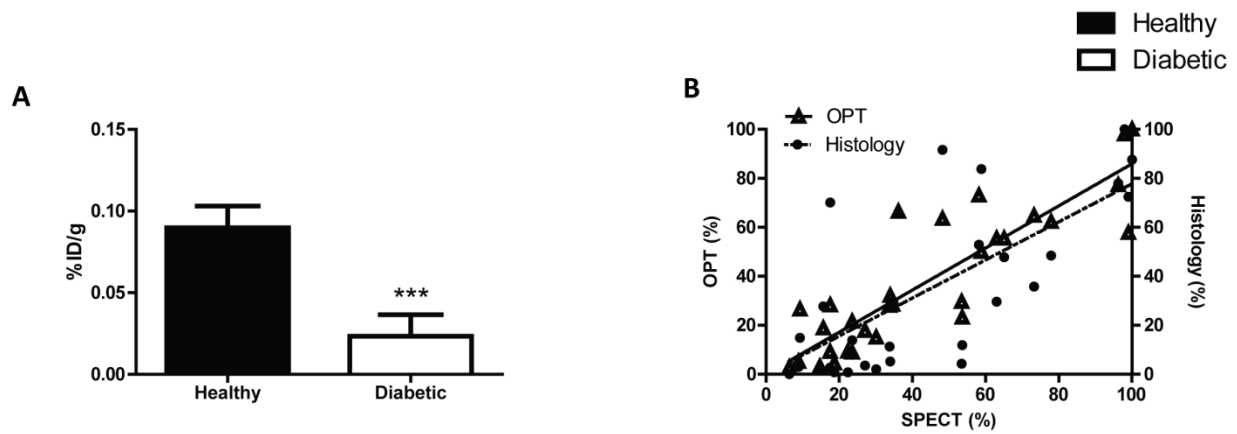
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**Figure S1. Linear correlations between  $^{111}\text{In}$ -exendin-3 uptake and BCM.** **A**, Correlation of SPECT data (kBq) with gamma-counter analysis (%ID, percentage of injected dose) of separate pancreatic lobes ( $r^2=0.85$ ,  $p=5.55 \times 10^{-12}$ ) ( $n=15$ ). **B and C**, graphs showing total pancreatic uptake of  $^{111}\text{In}$ -exendin-3 (SPECT) plotted against total BCM (histology) ( $r^2=0.53$ ,  $p=0.017$ ) and total pancreatic  $\beta$ -cell volume (OPT) ( $r^2=0.81$ ,  $p=0.0004$ ) respectively ( $n=5$ ).



**Figure S2. Alloxan treated rats display a significantly reduced uptake of  $^{111}\text{In}$ -exendin-3.** **A**, The Graph illustrates the pancreatic uptake of radioactivity normalized by the weight of the pancreas (%ID/g, percentage of injected dose per gram) of healthy control animals and alloxan treated animals as measured by gamma-counter. **B**, Two-way dependent correlation coefficients comparison (SPECT-OPT  $r^2=0.77$ , SPECT-histology  $r^2=0.52$ ;  $n=30$ ) reveals significant differences between SPECT-OPT and SPECT-histology correlations ( $p<0.001$ ). ANCOVA analysis reveals no significant differences between SPECT-OPT and SPECT-histology linear regression slopes ( $p=0.82$ ). Data are expressed in relation to (%) the highest BCM value measured in a lobe by the corresponding modality.